

Refine Search

Search Results -

Term	Documents
5.CLM..PGPB,USPT.	0
(L5.CLM.).PGPB,USPT.	0

Database:

US Pre-Grant Publication Full-Text Database
 US Patents Full-Text Database
 US OCR Full-Text Database
 EPO Abstracts Database
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 Derwent World Patents Index
 IBM Technical Disclosure Bulletins

Search:

L6

 





Search History

DATE: Monday, April 03, 2006 [Printable Copy](#) [Create Case](#)

<u>Set</u>	<u>Name</u>	<u>Query</u>	<u>Hit Count</u>	<u>Set Name</u>
	side by side			result set
	DB=PGPB,USPT; PLUR=YES; OP=ADJ			
<u>L6</u>	L5.clm.		0	<u>L6</u>
<u>L5</u>	(tgf\$ or transforming adj growth)same (antibod\$)same (nephritis or glomerulonephritis or diabetes)		195	<u>L5</u>
<u>L4</u>	(tgf\$ or transforming adj growth)same (antibod\$) and (nephritis or glomerulonephritis or diabetes)		1672	<u>L4</u>
<u>L3</u>	(L1 or L2) and (tgf\$ or transforming adj growth)		21	<u>L3</u>
<u>L2</u>	ruoslahti.in.		99	<u>L2</u>
<u>L1</u>	border.in.		320	<u>L1</u>

END OF SEARCH HISTORY

Search Results - Record(s) 1 through 10 of 21 returned.

1. 20050124534. 27 Sep 04. 09 Jun 05. Methods for treating conditions associated with the accumulation of excess extracellular matrix. Noble, Nancy A., et al. 514/8; 424/145.1 424/94.64 514/381 514/423 514/44 A61K048/00 A61K038/48 A61K039/395 A61K031/4178 A61K031/401.

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3. 20040028682. 07 Aug 03. 12 Feb 04. Inhibiting transforming growth factor beta to prevent accumulation of extracellular matrix. Border, Wayne A., et al. 424/145.1; 514/18 A61K039/395.

4. 20030045476. 20 Jul 01. 06 Mar 03. Heart homing conjugates. Ruoslahti, Erkki, et al. 514/16; A61K038/08.

5. 20030032591. 21 Aug 01. 13 Feb 03. Inhibitors of cell regulatory factors and methods for preventing or reducing scarring. Ruoslahti, Erkki I., et al. 514/12; 514/54 A61K038/17 A61K031/728.

6. 6906026. 05 Jul 01; 14 Jun 05. Methods for treating conditions associated with the accumulation of excess extracellular matrix. Noble; Nancy A., et al. 514/2; 435/4 435/6 435/7.1 514/4 514/44. A61K048/00 A61K039/395 A61K038/00.

7. 6509314. 13 Jun 94; 21 Jan 03. Methods of preventing or reducing scarring with decorin or biglycan. Ruoslahti; Erkki I.; et al. 514/8; 514/2 530/350. A01N061/00 A61K038/17 .

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Term	Documents
TRANSFORMING	110260
TRANSFORMINGS	3
GROWTH	416460
GROWTHS	7043
TGF\$	0

Search Results - Record(s) 11 through 20 of 21 returned.

11. 6277812. 02 Jun 95; 21 Aug 01. Methods for inhibiting TGF-.beta. activity. Ruoslahti; Erkki I., et al. 514/2; 435/69.1 514/8 530/395. A01N061/00 .

12. 6046162. 02 Jun 95; 04 Apr 00. Suppression of cell proliferation by decorin. Ruoslahti; Erkki I., et al. 514/8; 514/2 530/350 530/395. A61K038/02 A61K038/16 A61K038/39 .

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15. 5824655. 15 Feb 95; 20 Oct 98. Anti-transforming growth factor-.beta. gene therapy. Border; Wayne A.. 514/44; 424/93.21 424/93.7 435/320.1 435/352 435/353 435/354 435/366 514/2. A01N043/04 .

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Term	Documents
TRANSFORMING	110260
TRANSFORMINGS	3
GROWTH	416460
GROWTHS	7043
TGF\$	0
TGF	15708

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21. 5453492. 28 Jul 93; 26 Sep 95. 60 kDa transforming growth factor-.beta.-binding protein and its use to detect or purify TGF-.beta.. Butzow; Ralf, et al. 530/413; 435/7.1 530/350 530/395 530/402. C07K014/435 C07K001/22 G01N033/58 .

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Term	Documents
TRANSFORMING	110260
TRANSFORMINGS	3
GROWTH	416460
GROWTHS	7043
TGF\$	0
TGF	15708
TGFA	246
TGFAABMB	3
TGFAB	2
TGFACIDS	1
TGFAIL	1
((L1 OR L2) AND (TGF\$ OR TRANSFORMING ADJ GROWTH)).PGPB,USPT.	21

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L3: Entry 15 of 21

File: USPT

Oct 20, 1998

US-PAT-NO: 5824655

DOCUMENT-IDENTIFIER: US 5824655 A

TITLE: Anti-transforming growth factor-.beta. gene therapy

DATE-ISSUED: October 20, 1998

INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
<u>Border</u> ; Wayne A.	Salt Lake City	UT		

US-CL-CURRENT: 514/44; 424/93.21, 424/93.7, 435/320.1, 435/352, 435/353, 435/354,
435/366, 514/2

CLAIMS:

I claim:

1. A method for reducing deleterious accumulation of TGF-.beta. induced extracellular matrix in tissue of a subject, said method comprising:

(a) introducing into the subject a heterologous nucleic acid encoding a TGF-.beta. inhibitory agent selected from the group consisting of a proteoglycan, a TGF-.beta. receptor, an anti-TGF-.beta. antibody, a fragment of the proteoglycan, a fragment of the TGF-.beta. receptor, and a fragment of the anti-TGF-.beta. antibody, where each agent has the TGF-.beta. inhibitory function of reducing deleterious accumulation of TGF-.beta. induced extracellular matrix; and

(b) expressing in cells in said subject the introduced nucleic acid encoding the TGF-.beta. inhibitory agent and secreting the TGF-.beta. inhibitory agent encoded by the introduced nucleic acid for a time and under conditions effective to produce an amount of the TGF-.beta. inhibitory agent effective to reduce the deleterious accumulation of extracellular matrix material in said tissue of a subject where deleterious accumulation of extracellular matrix material induced by TGF-.beta. occurs.

2. The method of claim 1, wherein said TGF-.beta. specific inhibitory agent is first introduced into a cell *ex vivo* to obtain a cell expressing the TGF-.beta. specific inhibitory agent and the cell expressing said agent is transplanted into said subject.

3. The method of claim 1, wherein said cell is selected from the group consisting of a skeletal muscle cell, a kidney cell, a lung cell, a liver cell and a skin cell.

4. The method of claim 1, wherein said deleterious accumulation of extracellular matrix is associated with a condition in the subject selected from the group consisting of glomerulonephritis, ARDs, cirrhosis of the

livers, fibrotic cancer, fibrosis of the lungs, post myocardial infarction, cardiac fibrosis, post-angioplasty restinosis, renal interstitial fibrosis, scarring or a diabetes-associated pathology.

5. The method of claim 1, wherein said tissue is selected from the group consisting of kidney, liver, lung and skin tissue.

6. A method for reducing the accumulation of extracellular matrix associated with glomerulonephritis in a subject comprising:

a) introducing into the subject a heterologous nucleic acid encoding a TGF-.beta. inhibitory agent selected from the group consisting of a proteoglycan, a TGF-.beta. receptor, an anti-TGF-.beta. antibody, a fragment of the proteoglycan, a fragment of the TGF-.beta. receptor, and a fragment of the anti-TGF-.beta. antibody, where each agent has the TGF-.beta. inhibitory function of reducing deleterious accumulation of TGF-.beta. induced extracellular matrix; and

(b) expressing in cells in said subject the introduced nucleic acid encoding the TGF-.beta. inhibitory agent and secreting the TGF-.beta. inhibitory agent encoded by the introduced nucleic acid for a time and under conditions effective to produce an amount of the TGF-.beta. inhibitory agent effective to reduce accumulation of extracellular matrix material induced by TGF-.beta. in the kidney and suppress glomerulonephritis.

7. The method of claim 1 or claim 6, wherein said TGF-.beta. specific inhibitory agent is selected from the group consisting of decorin, biglycan, fibromodulin, lumican, betaglycan and endoglin, and fragments thereof having TGF-.beta. inhibitory action.